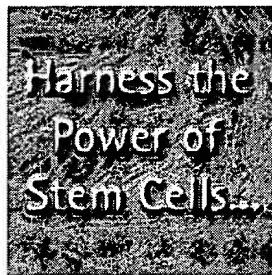
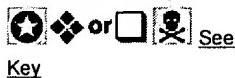


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Pale yellow solid. PROTECT FROM LIGHT. PACKAGED UNDER INERT GAS. A potent, analog of Cyclopamine (Cat. No. 239803) that specifically inhibits the Hedgehog (Hh) signal or lower toxicity ($IC_{50} = 20$ nM in Shh-LIGHT2 assay; 50 nM in p2^{Ptch-/-} cells; 500 nM in Sf9 cells). Binds to SmoA1 and promotes its exit from the endoplasmic reticulum. Suppresses induced pathway activity and SmoA1-induced reporter activity. *Purity:* ≥95% by TLC.

Ref.: Watkins, D.N., et al. 2003. *Nature* **422**, 313. Berman, D.M., et al. 2002. *Science* **297**, et al. 2002. *Proc. Natl. Acad. Sci. USA* **99**, 14071. Chen, J.K., et al. 2002. *Genes Dev.* **16**, Kamenetsky, M., et al. 2002. *J. Biol.* **1**, 10. Taipale, J., et al. 2000. *Nature* **406**, 1005.

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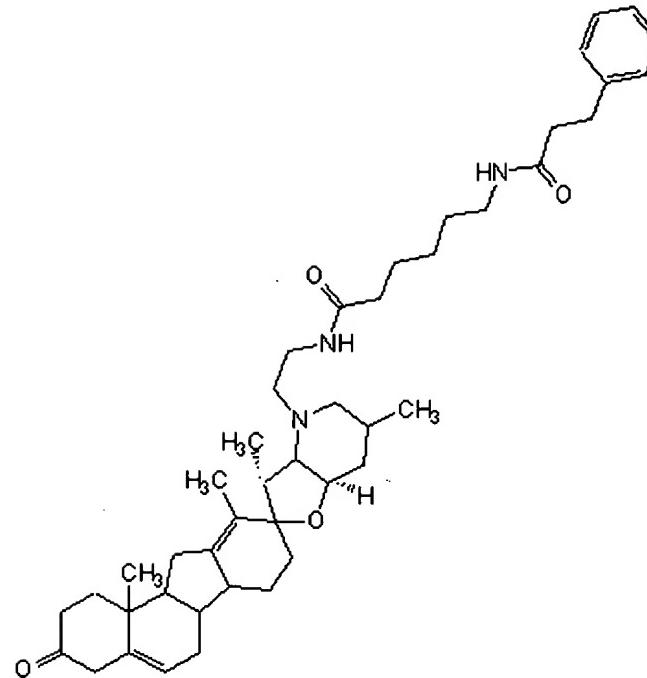
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100 µg	Y	<input type="text"/>	N/A

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Solubility

DMSO or MeOH

Molecular FormulaC44H63N3O4

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